

4. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste is soluble in water and slightly soluble in the wax.

5. (Amended) A granular pharmaceutical composition according to claim 1, wherein the wax has a melting point of 40-150°C.

6. (Amended) A granular pharmaceutical composition according to claim 1, wherein the wax is a member selected from the group consisting of hydrogenated oils, fats and oils of vegetable or animal origin, higher alcohols, polyethylene glycols, higher fatty acids, glycerin fatty acid esters, sucrose fatty acid esters, and combinations of two or more of these.

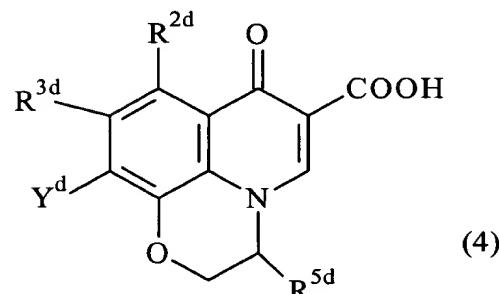
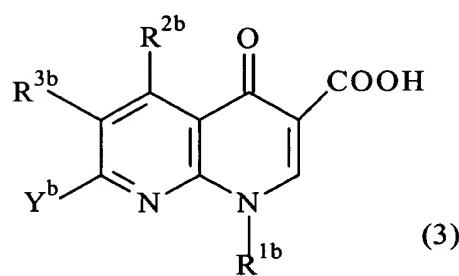
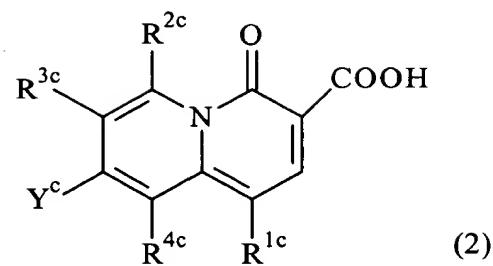
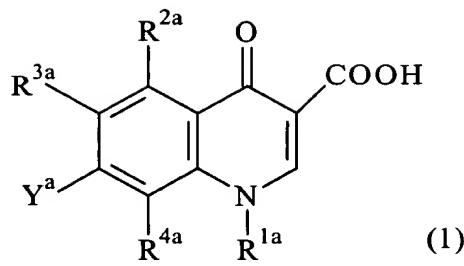
7. (Amended) A granular pharmaceutical composition according to claim 1, wherein the sugar alcohol is a member selected from the group consisting of erythritol, xylitol, sorbitol, maltitol, and combinations of two or more of these.

8. (Amended) A pharmaceutical composition according to claim 1, wherein the sugar alcohol has a heat of dissolution of not higher than -30 cal/g.

9. (Amended) A granular pharmaceutical composition according to claim 1, wherein the sugar alcohol is erythritol and/or xylitol.

10. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste is a drug selected from the group consisting of cetraxate hydrochloride, ecapapide, nefiracetam, talampicillin hydrochloride, indenolol

hydrochloride, hydralazine hydrochloride, chlorpromazine hydrochloride, tiaramide hydrochloride, berberine chloride, digitoxin, sulpyrine, azelastine hydrochloride, etilefrine hydrochloride, diltiazem hydrochloride, propranolol hydrochloride, chloramphenical, aminophyllin, erythromycin, clarithromycin, phenobarbital, calcium pantothenate, indeloxazine hydrochloride, aminoguanidine hydrochloride, bifemelane hydrochloride, 7 β -[2-(2-aminothiazol-4-yl)-2-(Z)-hydroxyiminoacetamido]-3-N,N-dimethylcarbamoyloxymethyl-3-cephem-carboxylic acid 1-(isopropoxycarbonyloxy)ethyl ester hydrochloride, (E)-3-(2-methoxy-3,6-dimethyl-1,4-benzoquinone-5-yl)-2-[5-(3-pyridyl)pentyl]-2-propenic acid, aminophylline, theophylline, diphenhydramine, metaclopramide, phenylbutazone, phenobarbital, ampicillin, cimetidine, famotidine, nizatidine, acetaminophen, epirizole, pyrazinamide, caffeine, ethionamide, carvedilol, ranitidine hydrochloride, roxatidine acetate hydrochloride, imipramine hydrochloride, ephedrine hydrochloride, diphenhydramine hydrochloride, tetracycline hydrochloride, doxycycline hydrochloride, naphazoline hydrochloride, noscapine hydrochloride, papaverine hydrochloride, dextrhomethorphan hydrobromide, timoepidium bromide, chlorphenilammonium maleate, alimemazine tartrate, pilsicainide hydrochloride, N-methylscopolamine methylsulfate, cinepazide maleate, arginine hydrochloride, histidine hydrochloride, lysine hydrochloride, lysine acetate, clopidogrel sulfate; crude drugs or extracts thereof; pyridonecarboxylic acid compounds represented by formulas (1) through (4) and salts thereof:



wherein each of R^{1a}, R^{1b}, and R^{1c} represent a C1-C6 linear or branched alkyl group which may have a substituent, a C3-C6 cyclic alkyl group which may have a substituent, an aryl group which may have a substituent, or a heteroaryl group which may have a substituent; each of R^{2a}, R^{2b}, R^{2c}, and R^{2d} represents a hydrogen atom or a C1-C6 linear or branched alkyl group which may have a substituent; or an amino group each of R^{3a}, R^{3b}, R^{3c}, and R^{3d} represents a hydrogen atom or a halogen atom; R^{4a} or R^{4c} represents a hydrogen atom, a halogen atom, a C1-C6 linear or branched alkyl group which may have substituent; or a C1-C6 linear or branched alkoxy group which may have a substituent; R^{5d} represents a hydrogen atom or a C1-C6 linear or branched alkyl group which may have a substituent; and each of Y^a, Y^b, Y^c, and Y^d represents a nitrogen-containing group.

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11. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste is ofloxacin.

12. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste is levofloxacin.

13. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste is clopidogrel sulfate.

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14. (Amended) A granular pharmaceutical composition according to claim 1, wherein the drug having a disagreeable taste and the wax are mixed at a ratio of 1:1 - 1:5 by weight, and the composition has a sugar alcohol content of at least 10% by weight.

15. (Amended) A granular pharmaceutical composition according to claim 1, which is produced by melting the wax with heat; dispersing or dissolving therein the drug having a disagreeable taste; subjecting the resultant mixture to primary granulation to thereby obtain a granulated product; and mixing the granulated product with the sugar alcohol, or subjecting the granulated product and the sugar alcohol to secondary granulation.

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19. (Amended) A pharmaceutical product for oral administration comprising the granular pharmaceutical composition claim 1.
